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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/081,400	02/20/2002	Michael Young	10275-041002	3033
31904	7590	04/09/2004	EXAMINER	
GTC BIOTHERAPEUTICS, INC. 175 CROSSING BOULEVARD, SUITE 410 FRAMINGHAM, MA 01702			WOITACH, JOSEPH T	
			ART UNIT	PAPER NUMBER
			1632	

DATE MAILED: 04/09/2004

Please find below and/or attached an Office communication concerning this application or proceeding.

DETAILED ACTION

This application filed February 20, 2002 is a division of 09/333,213 filed June 15, 1999, now US Patent 6,548,653, which claims benefit to provisional application 60/089343, filed June 15, 1998.

Applicants' preliminary amendment filed February 20, 2002, has been received and entered. Claims 27-45, 48-50 and 53-55 have been canceled. Claim 52 has been amended. Claims 1-26, 46, 47, 51 and 52 are pending and currently under examination.

Election/Restrictions

Restriction to one of the following inventions is required under 35 U.S.C. 121:

- I. Claims 1-26, 46 and 47, drawn to erythropoietin-serum albumin fusion protein, classified in class 530, subclass 399.
- II. Claims 51 and 52, drawn to an analog of erythropoietin, classified in class 530, subclass 402.

The inventions are distinct, each from the other because of the following reasons:

Inventions I and II are related as mutually exclusive species in an intermediate-final product relationship. Distinctness is proven for claims in this relationship if the intermediate product is useful to make other than the final product (MPEP § 806.04(b), 3rd paragraph), and the species are patentably distinct (MPEP § 806.04(h)). In the instant case, the intermediate product is deemed to be useful as analog of EPO itself without the addition of albumin and the inventions are deemed patentably distinct since there is nothing on this record to show them to be

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obvious variants. Moreover, each of the proteins encompassed by each restriction group are structurally and functionally different. Should applicant traverse on the ground that the species are not patentably distinct, applicant should submit evidence or identify such evidence now of record showing the species to be obvious variants or clearly admit on the record that this is the case. In either instance, if the examiner finds one of the inventions anticipated by the prior art, the evidence or admission may be used in a rejection under 35 U.S.C. 103(a) of the other invention.

Because these inventions are distinct for the reasons given above and have acquired a separate status in the art as shown by their different classification, and have acquired a separate status in the art because of their recognized divergent subject matter, restriction for examination purposes as indicated is proper.

Applicant is advised that the reply to this requirement to be complete must include an election of the invention to be examined even though the requirement be traversed (37 CFR 1.143).

Applicant is reminded that upon the cancellation of claims to a non-elected invention, the inventorship must be amended in compliance with 37 CFR 1.48(b) if one or more of the currently named inventors is no longer an inventor of at least one claim remaining in the application. Any amendment of inventorship must be accompanied by a request under 37 CFR 1.48(b) and by the fee required under 37 CFR 1.17(i).

In addition, for the sake of compact prosecution it is noted that the instant application contains sequences that are not identified by SEQ ID NOs., *for example* page 3 of the specification and claims 18, 19 and 23. The nucleotide sequence disclosure contained in this application does not comply with the requirements for such a disclosure as set forth in 37 C.F.R. 1.821 - 1.825. Applicant's attention is directed to the final rulemaking notice published at 55 FR 18230 (May 1, 1990), and 1114 OG 29 (May 15, 1990). If the effective filing date is on or after July 1, 1998, see the final rulemaking notice published at 63 FR 29620 (June 1, 1998) and 1211 OG 82 (June 23, 1998).

Appropriate correction is required.

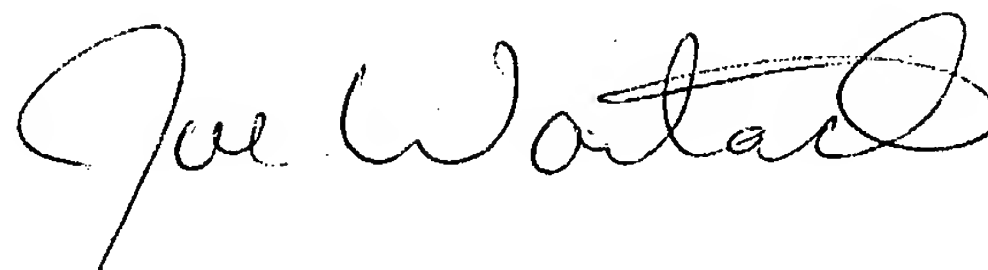
The absence of proper sequence listing did not preclude the restriction requirement however, **for a complete response to this office action, applicant must submit the required material for sequence compliance.**

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Joseph Woitach whose telephone number is (571) 272-0739.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Deborah Reynolds, can be reached at (571) 272-0734.

Any inquiry of a general nature or relating to the status of this application should be directed to the Group analyst Dianiece Jacobs whose telephone number is (571) 272-0532.

Joseph T. Woitach

A handwritten signature in black ink that reads "Joe Woitach". The signature is written in a cursive, flowing style with a large, prominent loop at the end of the last name.



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SERIAL NUMBER	FILING DATE	FIRST NAMED APPLICANT	ATTORNEY DOCKET NO.
10/081,400	2/20/2002	Young et al.	

EXAMINER	
Joseph Woitach	
ART UNIT	PAPER NUMBER
1632	04072004

Please find below a communication from the EXAMINER in charge of this application

This application contains sequence disclosures that are encompassed by the definitions for nucleotide and/or amino acid sequences set forth in 37 C.F.R. § 1.821(a)(1) and (a)(2). However, this application fails to comply with the requirements of 37 C.F.R. §§ 1.821-1.825 for the reason(s) set forth on the attached Notice To Comply With Requirements For Patent Applications Containing Nucleotide Sequence And/Or Amino Acid Sequence Disclosures. Applicant must comply with the requirements of the sequence rules (37 CFR 1.821 - 1.825) before the application can be examined under 35 U.S.C. §§ 131 and 132.

APPLICANT IS GIVEN **30 days** FROM THE DATE OF THIS LETTER WITHIN WHICH TO COMPLY WITH THE SEQUENCE RULES, 37 C.F.R. §§ 1.821-1.825. Failure to comply with these requirements will result in ABANDONMENT of the application under 37 C.F.R. § 1.821(g). Extensions of time may be obtained by filing a petition accompanied by the extension fee under the provisions of 37 C.F.R. § 1.136. **In no case may an applicant extend the period for response beyond the six month statutory period.** Applicant is requested to return a copy of the attached Notice to Comply with the response. Note that a reply to a notice to comply with the sequence rules should **not** be sent to the 20231 zip code address for the United States Patent and Trademark Office.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Joseph Woitach whose telephone number is (571)272-0739

Joseph T. Woitach

Joe Woitach
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